US ERA ARCHIVE DOCUMENT

DATA EVALUATION RECORD FISH LIFE-CYCLE TOXICITY TEST GUIDELINE 72-5

1. CHEMICAL: Disulfoton

Shaughnessey #: 032501

2. TEST MATERIAL: Di-Syston (non-radiolabeled) Purity: 98.5%

Radiolabeled Di-Syston-98.8% radiopurity

Radiolabeled Di-Syston Sulfone--99.3% Radiolabeled Di-Syston Sulfoxide--98.9%

3. CITATION:

Authors: Dionne, Emily

Title: Di-Syston - The chronic toxicity to the Sheepshead minnow (Cyprinodon variegatus)

during a full-life cycle exposure.

Study Completion Date: March 15, 1996

Laboratory: Springborn Laboratories, Inc., Environmental Sciences Division, 790 Main

St., Wareham, MA

Laboratory Report ID: 13507.0894.6110.592

Project ID Number: 107119

Sponsor: Bayer Corporation, Agriculture Division, P.O. Box 4913, Hawthorne Rd.,

Kansas City, MO 64120-0013

MRID No.: 439605-01

4. REVIEWED BY:

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Date: 10/5/97

5. APPROVED BY:

Daniel Rieder, Branch Chief, ERBIII, EFED, U.S.EPA

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Date: 6/18/0

6. STUDY PARAMETERS:

Test Species: Sheepshead minnow (Cyprinodon variegatus)

Age or Weight: embryos at test initiation

Duration of Test: 110 days Study Method: full life cycle test

Type of Concentration: nominal and mean measured

7. CONCLUSIONS:

This study appears to be scientifically sound but does not fulfill guideline requirements. Effects on F_o reproduction (fecundity), F_o growth, F_1 hatching success and F_o morphological abnormalities were seen at the lowest test level, so a NOEC was not



achieved for these parameters. The study does provide some information that is useful for a risk assessment, albeit with a high degree of uncertainty due to the deviations from guideline recommendations. Additional testing is required at levels low enough to produce a true NOEC.

8. ADEQUACY OF THE STUDY:

A. Classification: Supplemental

- B. Rationale: NOECs not achieved for several parameters, exposure tanks aerated, F₁ generation only exposed for half the recommended time
- C. Repairability: Additional testing at lower concentrations is required to achieve a true NOEC.

9. SUBMISSION PURPOSE: In support of reregistration of disulfoton

10. GUIDELINE DEVIATIONS:

Items not reported:

- 1) Did not report examination of embryos with scope or magnifying lens, as described in guidelines
- 2) Did not report percentage of healthy and fertile embryos from each spawn
- 3) Did not report time of day that embryos were counted and removed--guidelines suggest that it be done at the same time each day to minimize disruption of fish
- 4) Did not report the number of spawnings used to obtain embryos (should be at least three)
- 5) Did not report amount of time embryos soaked in dilution water prior to initiation of test (should be at least 2 hours)
- 6) Did not report whether live fungused embryos were removed and counted as dead
- 7) Did not report if lethargic or deformed juvenile fish were included in random selection
- 8) Did not report if survival was determined weekly -- only reports for days 28 and 57 post-hatch were included

Items which differ from guidelines:

1) Exposure solutions were aerated from day 55 through the end of the study--no data on daily DO levels were provided to show that this aeration was necessary, and no explanation of why the exposure tanks were aerated instead of aerating the dilution

water prior to input into the diluter system

- 2) The F₁ generation was only exposed for 4 weeks instead of the recommended 8 weeks
- 3) Males killed during spawning were replaced; guidelines recommend that neither males nor females killed in the spawning process should be replaced
- 4) No NOECs were achieved for several parameters: F_o reproduction (fecundity), F_o growth, F_1 hatching success and F_o morphological abnormalities
- 5) There were only two true replicates per treatment instead of the recommended four (there were four egg cups used, but these are not true replicates).

11.MATERIALS AND METHODS:

A. Biological System:

Guideline Criteria	Reported Information
Species: A freshwater fish species, preferably a fathead minnow (<i>Pimephales promelas</i> or an estuarine fish species, preferably a sheepshead minnow (Cyprinodon variegatus).	Estuarine speciesSheepshead minnow
Source and Acclimation of Fish 1. From wild population or Suitable laboratory culture 2.1. Sheepshead held in flowing 30°C seawater of >15% salinity for at least 2 wks. 2.2. Fathead 25°C and 16 hour/day day-light photoperiod (embryos will mature in 5 to 6 months under these conditions) 3. Neither species of fish or eggs should exhibit excess mortality.	 Embryos obtained from laboratory-held fish, which were purchased from Aquatic Biosystems as juveniles. They were held four and one-half months prior to being spawned. Held at 24-25°C, 31-32 % salinity for 14 days prior to spawning. Embryos were held at 25°C on a 12-hour photoperiod. These conditions were held throughout the test period. Pre-spawning fish exhibited a mortality rate of 2%.

Guideline Criteria	Reported Information
Eggs from Adult Fish Artificial inducement and natural spawning are the 2 methods for obtaining a sufficient number of eggs for a chronic exposure. 1. Artificial inducement (entails the stimulation of egg production by injection of human gonadotrophic hormone. Usually 10 \(\Perp \) s and 5 \(\Perp \) s should be used.) 2. Natural spawning (is possible with a few considerations for each fish species.)	 Natural spawning used. 2 males and 5 females from each test aquarium was placed in one section of the spawning chamber when they reached maturity (approx. 57 days post-hatch). They were held for 14 days.
3. Adult deaths during spawning should be noted; dead animals removed but not replaced.4. At termination of each spawning group, lengths and weights of individual fish are measured.	3. Females killed were not replaced; males were replaced to maximize egg fertilization success. 4. Each fish was measured, wet weighed, and internally examined.
Feeding	
1. Fry of both fish species should be fed equal portions of live brine shrimp nauplii at least 2x/day about 6 hours apart for three wks (frozen nauplii are not to be used).	1. Parent and first generation larvae were fed live brine shrimp nauplii 3x daily.
2. <u>Juveniles</u> (4 wks posthatch) and adults can be fed 2x/day on equal portions of dry food (e.g., Tetramin [®] or Biorell) supplemented with frozen adult brine shrimp.	2. Juvenile and adult fish were fed Zeigler Prime Flakes and frozen brine shrimp 2x daily.
3. Each batch of food should be checked for pesticides and metals.	3. Samples from each food source were analyzed periodically for the presence of pesticides, PCBs and toxic metals.
Embryo Removal	
1. Daily record numbers and egg fertility.	1. Eggs from the 28 brood stock groups were pooled and a representative sample of 200 were microscopically examined to estimate the percentage of successfully fertilized eggs. This method indicated that 76% were fertilized.

Guideline Criteria	Reported Information
2. Examined all embryos daily with a dissecting scope or magnifying viewer to remove empty shells and opaque, or abnormal embryos.	2. Embryos in cups were counted daily, and dead embryos were removed. There is no indication that the embryos were examined daily as described at left.
3. If <50% of the embryos from a spawn appear to be healthy and fertile, all embryos from that spawn should be discarded.	3. This information was not reported.
4. Embryos should be removed at a fixed time each day so spawning activity is not disturbed unnecessarily.	4. Dead embryos were removed daily, but time information was not reported.
Embryo Exposure (Four-Five Days)	
 The life-cycle chronic toxicity test must begin with embryos from at least 3 separate spawnings that are ≤24 hours old and have soaked in dilution water for at least 2 hours. 	 Information regarding number of spawns was not reported-there were 28 groups of 5 females and 2 males each of brood stock. Embryos were ≤ 24 hours old at initiation. Information not reported.
4. Testing begins by randomly distributing 50 embryos to each of the 4 replicate larval growth chambers.	4. 50 embryos impartially placed into 2 cups per replicate, 2 true replicates per treatment.
5. 10 embryos are transferred with a large bore eye dropper to successive incubation cups which are standing in dilution water. This is repeated until 50 embryos are in each cup. The incubation cups are then distributed to each replicate larval chamber.	5. Embryos were distributed to cups 5 at a time until each cup contained 50.

Guideline Criteria	Reported Information
6. 6.1 Survival of embryos, 6.2 time required to hatch, 6.3 hatching success, 6.4 and survival of fry for 4 wks are determined and recorded. 6.5.1 Dead embryos usually turn opaque and must be counted and 6.5.2 removed each day until hatching is complete. 6.6 Live fungused embryos must be removed daily 6.7. and counted as dead.	6. 6.1 Live embyros counted daily 6.2 reported for F ₁ only 6.3 percent hatching success reported for each replicate aquarium 6.4 28-day survival reported 6.5.1 dead counted 6.5.2 dead removed daily 6.6 Information not reported 6.7 Information not reported
Larval-Juvenile Exposure (Eight Weeks)	
 After hatching, each group of larvae is randomly reduced to 25, and released in replicate larval growth chambers. This random selection must include any fish that are lethargic or deformed. Survival should be determined in each replicate growth chamber at least once a week. Survival during this period is determined by counting the number of live fish, because dead larvae deteriorate rapidly. 	 Reduced to 25 from combined fish from 2 growth chambers Not reported Frequency of determination not reported; dead were removed daily so live were presumably counted at this time. Live fish counts used for survival
2. At 4 and 8 wks after hatching, total lengths (mm) of all fish must be recorded. 3. The amount of food given to the control and treated fish must be kept constant between exposures. Juvenile-Adult Exposure (32-40 wks)	2. Length and wet weight recorded at day 28; length recorded at day 57 also. Percent survival reported at days 28 and 57 3. Food amount not reported; larval fish were fed live brine shrimp nauplii three times daily; juvenile and adult fish were fed flake food and frozen brine shrimp daily. Food was withheld 24 hours before weight determination.

Guideline Criteria	Reported Information
 All fish are transferred to the adult spawning tank (same concentration) 8 wks after hatching. Each tank should have 25 randomly selected fish (deformed fish included). 	 Placed in spawning tank on day 57 post-hatch. 25 per tank; no report of deformed fish being included.
 3. When secondary sexual characteristics are well-developed, fathead minnow (20-24 week post hatch). Mature fish should be placed in spawning tank, separate from undeveloped fish. 4. The spawning tank will be divided into 4 individual spawning chambers with appropriate spawning substrates. 5. 4 ♂s and 4 ♀s are randomly chosen and assigned to spawning chambers. 6. Substrates are examined daily and embryos removed, counted, and recorded separately for each pair. 	 Majority had reached maturityall were placed in spawning tank on day 57. 2 separate spawning groups per replicate, 2 replicates per treatment. No description of substrates was provided. 2 males and 5 females per spawning chamber Spawns removed and counted daily
7. The adult exposure (fathead minnow) should be terminated when, during the decreasing day-length photoperiod, a 1-wk period passes in which no spawning occurs. 8. Testing using sheepshead minnows should terminate after spawning is observed for 2 wks because this fish spawns readily and almost daily unless immature or affected by a pollutant.	7. N/A 8. Testing terminated at 110 days post-hatch. Spawning occurred for 2 weeks.
Second Generation Embryo Exposure (4-5 days)	

Guideline Criteria	Reported Information
1. 50 embryos from each conc. level are randomly selected and transferred to incubation cups for hatch. 2. Those embryos not selected are discarded.	 50 embryos were selected from 2 groups of 50 per replicate from 2 different spawns, if possible Ones not used were discarded
Second Generation Larval-Juvenile Exposure (4-8 wks)	
 8 wk exposure begins with the release of 2 groups of 25 larvae in replicate growth chambers. These larvae should have been produced from different breeding pairs in each spawning tank. Selection of each group should be from early spawnings. 	 Exposure only lasted 28 days post-hatch From 2 spawns, if possible Embryos incubated from day of spawning
 4. Each group of 2nd generation fish is terminated 8 wks after hatching. 5. Fish are blotted, weighed, and measured before being discarded. 	4. Terminated at 4 weeks after hatching 5. Individual fish were weighed and measured

<u>Comments:</u> Second generation exposure of 4 weeks instead of 8 is not in agreement with the guideline recommendations.

B. Physical System:

Guideline Criteria	Reported Information
Test Water:	
Sheepshead Minnow 1. May be natural (sterilized and filtered) or a commercial mixture; 2. Natural seawater should have weekly range of salinity less than 6%, monthly pH range less than 0.8 pH units; 3. Salinity should be ≥ 15 parts per thousand; 4. Water must be free of pollutants.	 Natural filtered seawater Salinity ranged from 31-32 °/_∞ and pH ranged from 7.5-8.1 throughout the test period 31-32 parts per thousand salinity Analyzed routinelyfree of pollutants
Fathead Minnow	
1. Test water from well or spring which is not polluted 2. Sterilized and tested for pollutants 3. Hardness of 40 to 48 mg/L as CaCO ₃ and pH of 7.2 to 7.6 4. Reconstituted water can be used	N/A
Test Temperature: 1. For fathead minnow 25°C and should not remain outside the range of 24 to 26°C for more than 48 hours; 2. For sheepshead minnow, 30°C is recommended.	1. N/A 2. Test temperature was 25± 1°C

Guideline Criteria	Reported Information
Photoperiod:	1. N/A
1. Simulate wavelength spectra of sunlight	2. 12-h L/12-h D
Intensity 10 to 100 lumens at water surface.	3. N/A
 Sheepshead 12-hour light/12-hours dark Fathead dawn-to-dusk at Evansville, IN as of Dec. 1st 	
Dosing Apparatus: 1. Intermittent flow proportional diluters or continuous flow serial	Intermittent flow proportional diluter
diluters should be used. 2. A minimum of 5 toxicant	2. Yes- 5
concentrations	3. Yes- 0.5
3. with a dilution factor not greater than 0.5and4. 1 control should be used.	4. Negative control and solvent control were used
Toxicant Mixing: 1. Mixing chamber is recommended	1. Mixing chamber was used
but not required; 2. Aeration should not be used for mixing;	2. Not aerated for mixing
3. It must be demonstrated that the test solution is completely mixed	3. Yes
before intro. into the test system; 4. Flow splitting accuracy must be	
within 10% and periodically checked.	1 Vas ±5%
Test Vessels: All glass or glass with a plastic or stainless steel frame.	4. Yes, ±5% Yes, glass

Guideline Criteria	Reported Information
Fathead 1. Adult spawning tanks should measure 30.5 x 30.5 x 91.4 cm or 30.5 x 30.5 x 61 cm long with screened-off or separate larval tank. 2. Each larval section is divided in half allowing for two larval growth chambers for each adult spawning tank. 3. Larval chambers should be designed with glass bottoms and drains that allow water to be drawn down to 3 cm. 4.1. Test water must be delivered separately to each adult tank and larval section, 4.2 with one-third of the water volume going to the latter. 5. Test water depth in adult tanks and larval chambers should be a minimum of 15 cm.	N/A
Sheepshead 1. Tanks 45 x 90 x 26 cm with water depth of 19 cm recommended. 2. Larval chamber design and test water divided are the same as described for the fathead minnow.	 No, 60 x 30 x 30 cm, volume of 27L, depth of water not reported Yes5 cm glass jars with nitrex screen; larval chambers 30 x 13 x 25 cm glass with nitrex screen
Embryo and Fry Chambers: 1. 120 ml glass jars with bottoms replaced with 40 mesh stainless steel or nylon screen. 2. Chambers can be oscillated vertically (2.5 to 4.0 cm) (rocker arm apparatus, 2 rpm motor) or placed in separate chambers with self-starting siphons.	1. Yes 2. Not reported

Guideline Criteria	Reported Information
Flow Rate: 1. Flow rates to larval cups should provide 90% replacement in 8-12 hours. 2. Flow rate must maintain DO at above 75% of saturation and maintain the toxicant level (cannot drop below 20% with fish in the	 Yes-7.4 volume replacements in a 24 hour period (90% replacement in 7.5 hours) Nobegan aeration at day 55 to maintain DO level
Aeration: 1. Dilution water should be aerated to insure DO concentration at or near 100% saturation. 2. Test tanks and embryo chambers should not be aerated.	Nothey aerated individual exposure aquaria instead of the dilution water 2. Nothey were aerated

C. Chemical System:

Guideline Criteria	Reported Information
Concentrations: 1.1 Minimum of 5 concentrations and a control,	1. Five conc. and 2 replicates
1.2 all replicated, plus solvent control if appropriate.2 Toxicant conc. must be measured in	1.2. Yes
one tank at each toxicant level every week. 3 One concentration must adversely affect a life stage and one concentration	2. Yes3. No, all levels were affected for the F₀
must not affect any life stage.	abnormalities, F_0 growth, F_0 fecundity, and F_1 hatching success.

Guideline Criteria	Reported Information
Other Variables: 1. DO must be measured at each conc. at least once a week; 2. Freshwater parameters in a control and one conc. must be analyzed once a week for pH, alkalinity, hardness, and conductance 3. Natural seawater must maintain a constant salinity and not fluctuate more than 6% weekly; monthly pH range < 0.8 pH units.	1. Yes, daily 2. N/A 3. Yes
Solvents: Should not exceed 0.1 ml/L in a flow-through system. Following solvents are acceptable: dimethylformamide, triethylene glycol, methanol, acetone, ethanol.	Solvent: acetone, 1.8 μL/L

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12. REPORTED RESULTS:

Reported Statistical Results for Biological Endpoints:

Guideline Criteria	Reported Information
Data Endpoints must include:	
 survival of F₀ and F₁ embryos, time required to hatch, hatching success, and survival of fry 	1. Yes 2. Yes (F ₁ only) 3. Yes 4. Yes
 survival of F₀ fish during larval-juvenile exposure period 1 at 4 and 8 weeks after hatching, 2. total lengths of fish 1 at 8 weeks after hatching of F₁ fish, 2. weights and lengths are recorded 	1. Yes 2. Yes
 incidence of pathological or histological effects observations of other effects or clinical signs 	 Yesat 4 weeks after hatching Yes Yesphysical deformations, status of gonads lethargy, etc.

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Table 1: Hatching Success, Larval Survival and Growth of the F_o Sheepshead Minnow after 28 and 57 days post-hatch exposure to Di-Syston

			Day 28			Day 57		
Mean Measured Concenration (μg/L)	Percent Hatching* [% reduction from pooled control]	Percent Larval Survival [% reduction from pooled control]	Length ^b in mm (std. dev.) [% reduction vs pooled control]	Wet Weight ^b in g (std. dev.) [% reduction vs pooled control]	Percent Larval Survival° [% reduction vs pooled control]	Length ^b in mm (std. dev.) [% reduction vs pooled control]		
Control A B Mean	70 61 66	96 96 96	27.3 (2.0) 26.5 (1.7) 26.9 (1.9)	0.30 (0.05) 0.28 (0.05) 0.29 (0.05)	100 100 100	40.0 (4.9) 39.8 (4.2) 39.9 (4.5)		
Solvent A Control B Mean	70 77 74	94 96 95	27.3 (1.8) 27.2 (1.7) 27.3 (1.7)	0.32 (0.05) 0.31 (0.04) 0.32 (0.04)	100 100 100	41.1 (4.5) 41.0 (4.0) 41.0 (4.2)		
Pooled Control	70	96	27.1 (1.8)	0.30 (0.05)	100	not usedf		
2.9 A B Mean	64 65 65 [7%]	94 96 95 [1%]	27.0 (2.0) 26.6 (2.2) 26.8(2.1) [1.1%] °	0.32 (0.05) 0.30 (0.05) 0.31 (0.05) [0%]°	100 96 98 [2%]	38.4 (4.7) 38.7 (2.9) 38.6 (3.8) [6%] ^g		
5.7 A B Mean	59 66 63 [10%]	96 80 88 [8%] ^d	25.4 (1.9) 25.8 (1.5) 25.5 (1.7) [6%] ^h	0.29 (0.06) 0.32 (0.05) 0.30 (0.05) [0%] ^h	100 96 98 [2%]	37.5 (3.1) 36.5 (3.8) 37.1 (3.5) [10%] ²		
11 A B Mean	70 75 73 [0%]	82 92 87 [9%] ^d	24.4 (2.8) 23.8 (2.3) 24.1 (2.6) [11%] ^h	0.31 (0.09) 0.27 (0.06) 0.29 (0.07) [3%] ^h	100 92 96 [4%]	35.0 (3.4) 33.9 (5.4) 34.5 (4.5) [16%] [§]		
24 A B Mean	64 63 64 [9%]	78 72 75 [22%] ^d	22.2 (2.9) 21.8 (2.6) 22.0 (2.8) [19%] ^h	0.26 (0.09) 0.21 (0.06) 0.24 (0.08) [20%] ^h	96 100 98 [2%]	32.0 (5.7) 31.8 (4.3) 31.9 (5.0) [22%] ⁸		
47 A B Mean	70 68 69 [1%]	64 74 69 [28%] ^d	16.7 (4.2) 18.4 (3.6) 17.6 (3.9) [35%] ^h	0.06 (0.07) 0.012 (0.09) 0.10 (0.09) [67%] ^h	68 72 70 [30%] ^d	20.7 (4.2) 22.0 (4.6) 21.4 (4.4) [48%] ^h		

^aPercentage is based on the total number of eggs included in each replicate aquarium. A sub-sample viability determination indicated approximately 76% of these eggs were viable,

^bMeasurement presented as mean + standard deviation.

Percentage is based on the survival among larval groups of 25 which were established at day 28 post-hatch thinning of larvae

Table 2: Survival and Growth of Sheepshead Minnow at Test Termination (Day 110 Post-Hatch).

		Mean Total Len dev.) [% reduction control]	gth ^b in mm (std. on vs pooled	Mean Wet Weddev.) [% reduction	
Mean Measured Concentration (μg/L)	Percent Survival* [% reduction vs solvent control]	Male	Female	Male	Female
Control A B Mean	100 92 96	48.1 (4.6) 47.6 (3.9) 47.8 (4.2)	43.6 (3.4) 42.8 (1.7) 42.7 (2.7)	2.04 (0.48)	1.47 (0.24)
Solvent A Control B Mean	88 72 80	47.6 (3.7) 48.8 (3.9) 48.3 (3.7)	44.3 (2.7) 42.6 (2.5) 43.6 (2.7)	2.05 (0.48) 2.26 (0.67) 2.17 (0.58)	1.62 (0.35) 1.48 (0.32) 1.56 (0.34)
Pooled Control	e	48.0 (4.0)	43.2 (2.7)	2.10 (0.51)	1.54 (0.31)
2.9 A B Mean	100 92 96 [0%]	46.5 (3.3) 47.4 (2.8) 46.9 (3.1) [2%]	42.0 (2.2) 42.4 (2.9) 42.2 (2.6) [2%]	2.03 (0.43) 2.14 (0.40) 2.08 (0.41) [1%]	1.49 (0.29) 1.43 (0.32) 1.46 (0.30) [5%]
5.7 A B Mean	⁻ 96 100 98 [0%]	44.0 (4.3) 42.8 (3.1) 43.6 (3.8) [9%] ^f	41.1 (1.8) 40.3 (3.0) 40.6 (2.6) [6%] ^f	1.86 (0.54) 1.77 (0.41) 1.82 (0.49) [13%] ^f	1.37 (0.22) 1.33 (0.26) 1.35 (0.24) [12%]
11 A B Mean	100 84 92 [0%]	42.9 (4.4) 43.2 (2.9) 43.0 (3.8) [10%] ^f	37.9 (2.9) 34.9 (5.3) 36.4 (4.5) [16%] ^f	2.00 (0.70) 1.99 (0.40) 1.99 (0.59) [6%]	1.24 (0.28) 0.97 (0.41) 1.10 (0.37) [29%] ^f
24 A B Mean	96 92 94 [0%]	39.9 (5.2) 40.9 (3.2) 40.1 (4.4) [16%] ^f	33.3 (4.6) 34.1 (4.6) 33.8 (4.5) [22%] ^f	1.52 (0.62) 1.55 (0.40) 1.54 (0.54) [27%] ^f	0.79 (0.42) 0.83 (0.32) 0.81 (0.36) [47%] ^f

dSignificantly different (p \leq 0.05) at this treatment level compared to pooled control data.

[&]quot;No effect concentration based on t-test of treatment data vs pooled control data

Comparison made to solvent control data

⁵ Significantly different (p \leq 0.05) compared to solvent control data. The reduction at the 2.9 μ g/L treatment level was small (6%) and was considered biologically insignificant.

^hData not included in statistical comparison due to a survival effect at this treatment level.

47 A B	56 40	35.9 (5.2) 34.5 (1.1)	26.0 (3.1) 29.6 (5.7)	0.99 (0.42) 0.86 (0.19)	0.31 (0.12) 0.53 (0.36)
Mean	48	35.5 (1.5)	27.8 (4.8)	0.95 (0.36)	0.42 (0.28)
	[40%]°	[26%] ^s	[36%] ^g	[55%] ^g	[73%]8

Percent survival of fish between days 57 post-hatch and test termination.

Table 3: Egg Production of Fo Sheepshead Minnows Exposed to Di-Syston

		Mean # Eggs/	/Female/Day ^b
Mean Measured Concentration (μg/L)	Total # Eggs Produceda	Replicate Means	Treatment ^o Mean [% reduction vs pooled control]
Control A B	4803 3468	34.3 26.2	30
Solvent Control A B	6597 4115	47.1 30.0	39
Pooled Control	4746		34
2.9	2935 3299	21.0 23.6	22 [35%]4
5.7	2169 2887	15.5 20.6	18 [47%]°
11	1802 1203	12.9 8.6	11 [68%]°
24	185 315	1.3 2.3	2.0 [94%]*
47	0		[100%]

^aBased on the production of 2 spawning groups (2 males, 5 females each) for 14 days.

bStandard deviation presented in parentheses.

Significantly different (p \leq 0.05) at this treatment level compared to the solvent control data.

dWeight data for the (A) replicate were inadvertently lost when electronically stored during the test termination procedure.

^cComparison made to solvent control data.

Significantly different (p \leq 0.05) at this treatment level compared to the pooled control using Dunnett's test

⁸Data not included in statistical comparison due to survival effect at this treatment level.

b# eggs/female/day was calculated with the number of females alive on each day of spawning.

Rounded to whole numbers.

dSignificantly reduced compared to the pooled controls using Williams' Test. A NOEC was calculated by linear regression for this endpoint.

[&]quot;Significantly reduced compared to the pooled controls using Dunnett's Test.

^fNo eggs produced at this test concentration.

Table 4: Time-to-Hatch for F_1 Sheepshead Minnow Embryos Exposed to Di-Syston

Number of Groups Hatching Following			
Mean Measured Concentration (μg/L)	5 days	6 days	7 days
Control	0 10 / 10	31 (91)*	3 (9)
Solvent Control	1 (3)	26 (76)°	7 (21)°
2.9	0	32 (94)	2 (6)
5.7	0	32 (94)	2 (6)
11	1 (6)	16 (89)	1 (6)
24	1 (14)	6 (86)	0
47°	<u></u>		

^aPercentage of total number is presented in parentheses.

Morphological and Behavioral Observations

Parental Generation

1) "Dorsal Hump" deformity observed in F_o fish at all treatment levels, but not in the control fish. This was caused by a spinal curvature on the vertical plane and was considered lordosis. The numbers/percentage of affected fish is presented below:

Table 5: Numbers and Percentages of Fish Exhibiting "Dorsal Hump" in the F_0 Generation Treated With Di-Syston.

Mean Measured Concentration (μg/L)	Number of Fish	% of Total
control	0 1 52 40 52	0
solvent control	0	0
2.9	2	4
5.7	8	16
11	18	39
24	16	34
47	12	50

^bNo spawning occurred at this treatment level.

This percentage was lowered by the hatching time in the (B) replicate, where 53% of the embryo groups hatched in 6 days and 41% hatched in 7 days. In the (A) replicate, 100% hatched in 6 days.

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Lab's Conclusion: Dorsal hump deformity was treatment related and was seen at all treatment levels.

F1 Generation

1) No deformities/behavioral abnormalities seen

Other Observations

Raw data included? Yes

Statistical Results:

Statistical Method: Continuous data (growth and reproduction endpoints): control and solvent control were compared with a 2-tailed T-test. If no significant differences were seen, the two controls were pooled, and the treatments were compared to the pooled control using Dunnet's and William's Tests. If these tests were not suitable due to failure to meet the assumptions required, a Kruskal-Wallis (non-parametric) test was used.

Survival and hatching success data: a chi-square test was used to compare the control and solvent control data. If no significant difference was observed, the controls were pooled for analysis. Treatment results were compared to the pooled control (or solvent control if controls were not pooled) using either Fisher's Exact Test or Chi-Square.

NOEC: $0.96^1 \mu g/L$ LOEC: $2.9 \mu g/L$

MATC: 1.7 μ g/L

Most sensitive endpoint: F₀ fecundity (eggs/female/day)

¹This NOEC was extrapolated using linear regression. Statistically significant effects for this parameter were observed at all levels tested.

<u>Comments:</u> F_0 survival, growth, fecundity and F_1 hatching success were all significantly affected at 2.9 μ g/L, the lowest level tested, as was the occurrence of the dorsal hump deformity. Therefore, no true NOECs were determined for these parameters.

13. Reviewer's Discussion:

Statistical Results

Statistical Method: Dunnett's and Williams Tests, comparison of treatment vs solvent control.

NOEC: $<2.9 \mu g/L$ LOEC: $2.9 \mu g/L$ MATC: Could not be determined due to no NOEC obtained.

Most Sensitive Endpoint: Survival, growth, physical abnormalities, and fecundity for parental generation (F_0) fish, as well as hatching success for F_1 fish, were all significantly affected at the lowest level tested.

Significant Items

The lack of a NOEC for these multiple parameters makes it extremely difficult to use this data with any degree of certainty in a risk assessment. The many other deviations from Guideline requirements in this study (e.g., aeration of exposure aquaria, short F_1 exposure, etc.) also make the use of the data from this study more questionable in a risk assessment.

Run on law data Conc. listed as nominal

TITLE: Disyston Fish Early Life--Length Data FILE: a:dsfelslg.dat TRANSFORM: NO TRANSFORMATION NUMBER

NUMBER OF GROUPS: 7

Fo	· ·

GRP	IDENTI	FICATION	REP	VALUE	TRANS VALUE
1	solvent		1000	34.0000	34.0000
1		*	2	36.0000	36.0000
1	solvent		3	39.0000	39.0000
1	solvent		4	46.0000	46.0000
1	solvent		4 5	42.0000	42.0000
1	solvent		6	35.0000	35.0000
1	solvent		7	44.0000	44.0000
1	solvent	control	8	51.0000	51.0000
1	solvent	control	9	41.0000	41.0000
1	solvent	control	10	44.0000	44.0000
1	solvent	in the second	.11	44.0000	44.0000
1	solvent		12	37.0000	37.0000
1	solvent		13	46.0000	46.0000
1	solvent		14	44.0000	44.0000
1	solvent		15	40.0000	40.0000
1	solvent		16	45.0000	45.0000
1 1	solvent solvent		17 18	43.0000 45.0000	43.0000 45.0000
1	solvent	'	19	43.0000	43.0000
1	solvent		20	39.0000	39.0000
1	solvent		21	34.0000	34.0000
1	solvent		22 V .	39.0000	39.0000
1	solvent		23	35.0000	35.0000
1	solvent		24	36.0000	36.0000
1	solvent	control	25	42.0000	42.0000
1	solvent	control	26	40.0000	40.0000
. 1	solvent		27	42.0000	42.0000
1	solvent		28	43.0000	43.0000
1	solvent		29	45.0000	45.0000
1		control_		39.0000	39.0000
1	solvent		31	39.0000	39.0000 44.0000
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	solvent		34	47.0000	47.0000
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1	solvent		36	46.0000	46.0000
1	solvent		37	39.0000	39.0000
1	solvent		38	36.0000	36.0000
1 1	solvent		39	46.0000	46.0000
1, 1	solvent	control	40	40.0000	40.0000
′ 1 ·	solvent		41	46.0000	46.0000
1 1	solvent		42	34.0000	34.0000
1	solvent		43	38.0000	38.0000
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2		control	4	42.0000	42.0000
2		control	5	40.0000	40.0000
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2	contr	ol 17'	32.0000	32.0000	
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2	contr	ol 24'	40.0000	40.0000	
	contr		36.0000	36.0000	
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2	contr		40.0000	40.0000	
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2	contr	ol_ 46	39.0000	39.0000	
2	contr		42.0000	42.0000	
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. 7	50	12	27.0000	27.0000
7	50	13	26.0000	26.0000
7	50	14	27.0000	27.0000
7	50	15	22.0000	22.0000
7	50	16	16.0000	16.0000
7		17	16.0000	16.0000
7	50	18	20.0000	20.0000
7		19	30.0000	30.0000
	50	20	24.0000	24.0000
7	50	21	14.0000	14.0000
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	· · · · · · · · · · · · · · · · · · ·	23	22.0000	22.0000
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7	50	25	22.0000	22.0000
. 7	50	26	23.0000	23.0000
· 7	50	27	27.0000	27.0000
7	50	28	16.0000	16.0000
.7	50	29	25.0000	25.0000
7	50	30	23.0000	23.0000
7.7	50	31	27.0000	27.0000
7	50	32	24.0000	24.0000
7.	, , , , , ,	33	13.0000	13.0000
7	50	34	16.0000	16.0000
7	50	35	23.0000	23.0000
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Disyston Fish Early Life--Length Data File: a:dsfelslg.dat Transform: NO TRANSFORMATION

#### SUMMARY STATISTICS ON TRANSFORMED DATA TABLE 1 of 2

GRP	IDENTIFICATION	N	MIN	MAX	MEAN
1 2 3 4 5 6 7	solvent control control 3.1 6.3 13 25 50	44 50 47 48 45 45	34.000 30.000 30.000 28.000 19.000 20.000 13.000	51.000 54.000 51.000 45.000 41.000 40.000 30.000	41.045 39.880 38.574 37.063 34.467 31.889 21.371

Disyston Fish Early Life--Length Data
File: a:dsfelslg.dat Transform: NO TRANSFORMATION

#### SUMMARY STATISTICS ON TRANSFORMED DATA TABLE 2 of 2

GRP	IDENTI	FICATION	VARIANCE	SD	SEM
1	solvent	control	18.044	4.248	0.640
2		control	20.353	4.511	0.638
3		3.1	14.815	3.849	0.561
4		6.3	12.017	3.467	0.500
5		13	20.209	4.495	0.670
6		25	24.646	4.965	0.740

19.299

Disyston Fish Early Life--Length Data
File: a:dsfelslg.dat Transform: NO TRANSFORMATION

# ANOVA TABLE

SOURCE DF		SS	MS	F
Between 6	1.	10493.789	1748.965	95.042
Within (Error) 307		5649.307	18.402	
Total 313		16143.096		

Critical F value = 2.18 (0.05,6,120)

Since F > Critical F REJECT Ho: All groups equal

Disyston Fish Early Life--Length Data

File: a:dsfelslg.dat Transform: NO TRANSFORMATION

1	BONFERRONI T-TEST -	TABLE 1 OF 2	Ho:Contro	l <treatm< th=""><th>ent</th></treatm<>	ent
GROUP	IDENTIFICATION	TRANSFORMED MEAN	MEAN CALCULATED IN ORIGINAL UNITS	T STAT	SIG
1 2 3 4 5 6 7	all S.d.	いきょう こうこうこうじょ しょくじ	41.045 39.880 38.574 37.063 34.467 31.889 21.371	1.314 2.746 4.449 7.234 10.068 20.249	* * * *

Bonferroni T table value = 2.43 (1 Tailed Value, P=0.05, df=120,6)

Disyston Fish Early Life--Length Data

File: a:dsfelslg.dat Transform: NO TRANSFORMATION

	BONFERRONI T-TEST - TABLE	2 OF 2	Ho:Control <treatment< th=""></treatment<>
GROUP	NUM OF IDENTIFICATION REPS	Minimum Sig Diff (IN ORIG. UNITS)	
1 2 3 4 5 6 7	solvent control 44 control 50 3.1 47 6.3 48 13 45 25 45 50 35	2.154 2.186 2.175 2.209 2.209 2.360	5.2 1.165 5.3 2.471 5.3 3.983 5.4 6.579 5.4 9.157 5.7 19.674

Disyston Fish Early Life--Length Data File: a:dsfelslg.dat Transform: Transform: NO TRANSFORMATION

WILLIAMS TE	ST (Isotonic	regression	model)	TABLE	1	OF	2
-------------	--------------	------------	--------	-------	---	----	---

GROUP IDENTIFICAT	rion	N	ORIGINAL MEAN	TRANSFORMED MEAN	ISOTONIZED MEAN
1 solvent 2 3 4 5 6 7	control	44 50 47 48 45 45	41.045 39.880 38.574 37.063 34.467 31.889 21.371	41.045 39.880 38.574 37.063 34.467 31.889 21.371	41.045 39.880 38.574 37.063 34.467 31.889 21.371

Disyston Fish Early Life--Length Data
File: a:dsfelslg.dat Transform: NO TRANSFORMATION

WILLIAMS	TEST (Isotonic	regression	model)	TABLE 2 OF	' <b>2</b>
IDENTIFICATION	ISOTONIZED MEAN	CALC. WILLIAMS	SIG P=.05	TABLE WILLIAMS	DEGREES OF FREEDOM
m.	3.1 2.9 38.574 6.3 5.7 37.063 13 11 34.467 25 24 31.889 50 47 21.371	1.314 2.746 4.449 7.234 10.068 20.249	* * *	1.66 1.73 1.75 1.77 1.77	k= 1, v=307 k= 2, v=307 k= 3, v=307 k= 4, v=307 k= 5, v=307 k= 6, v=307

Note: df used for table values are approximate when v > 20.